

A randomised double blind placebo controlled trial of the addition of metformin to optimal treatment in patients with chronic heart failure. Assessment by regional haemodynamics neurohumoral activation and symptomatic well being

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 27/10/2015	Condition category Circulatory System	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
N0192080686

Study information

Scientific Title

A randomised double blind placebo controlled trial of the addition of metformin to optimal treatment in patients with chronic heart failure. Assessment by regional haemodynamics neurohumoral activation and symptomatic well being

Study objectives

To determine whether metformin is of benefit in the treatment of patients with congestive heart failure.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Cardiovascular: Congestive heart failure (CHF)

Interventions

Randomised controlled trial.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Metformin

Primary outcome(s)

1. Regional haemodynamics
2. Neurohumoral
3. Symptomatic well being

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/08/2004

Eligibility

Key inclusion criteria

Total number of subjects = 20

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

15/02/2000

Date of final enrolment

31/08/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Nottingham University Hospitals NHS Trust

Nottingham

United Kingdom

NG7 2UH

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Nottingham University Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration